

EXHIBIT E6

SUPERIOR COURT OF NEW JERSEY
LAW DIVISION: MIDDLESEX COUNTY

RICARDO RIMONDI and
PILAR RIMONDI,

Plaintiffs,

vs.

BASF CATALYSTS LLC, et al.,

Defendants.

DOCKET NO.

MID-L-2912-17

JOANNA RUMAN and
JACENTY RUMAN,

Plaintiffs,

vs.

BASF CATALYSTS LLC, et al.,

Defendants.

DOCKET NO.

MID-L-2919-17

DEPOSITION OF

WILLIAM E. LONGO, PhD

January 7, 2019

10:30 a.m.

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Johns Creek, Georgia

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14:07:07 **1 nonsensical definition to be asbestiform. So we know**
 14:07:10 **2 the bundles are asbestiform, we have sources for**
 14:07:14 **3 that.**
 14:07:15 **4 Now we're dealing with the single fibers.**
 14:07:20 **5 I don't believe it's reasonable to say 10 percent of**
 14:07:24 **6 the single fibers from these mine sources are unknown**
 14:07:28 **7 when you have all these other bundles. I don't think**
 14:07:33 **8 it's reasonable to say that the average aspect ratio**
 14:07:36 **9 from ground talc on tremolite that we find in these**
 14:07:40 **10 samples are the same as what Blount found as well as**
 14:07:44 **11 Campbell found as well as Langer found, saying these**
 14:07:49 **12 are all asbestiform with that aspect ratio. So**
 14:07:54 **13 that's my opinion, that these are asbestiform.**
 14:07:57 **14 Q.** Okay. So if I asked you in 2016 what your
 14:08:03 **15** definition of asbestiform was, what would you have
 14:08:05 **16** said?
 14:08:07 **17 MR. HORN:** Object to form. Vague and
 14:08:11 **18** ambiguous.
 14:08:11 **19 THE WITNESS:** I don't know.
 14:08:12 **20 Q.** (By Mr. Ewald) If you have used the word
 14:08:17 **21** asbestiform in prior transcripts before talc
 14:08:20 **22** litigation -- I'll withdraw that.
 14:08:24 **23** The definition you gave today of
 14:08:29 **24** asbestiform, is that the definition that you have
 14:08:32 **25** used the entire time that you have been testifying in
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14:08:37 **1** cosmetic talc litigation?
 14:08:40 **2 A. I think what I've stated in the past is**
 14:08:43 **3 that for single fibers you can't tell if it's**
 14:08:48 **4 asbestiform or not based on the definition. But**
 14:08:53 **5 we're not looking just at a single fiber. We're**
 14:08:55 **6 looking at multiple fibers and bundles.**
 14:08:58 **7 So you have the geological definition of**
 14:09:02 **8 asbestiform, and then you just have is it fibrous,**
 14:09:06 **9 therefore it's asbestiform.**
 14:09:07 **10 Q.** Under the if it's fibrous, therefore it's
 14:09:12 **11** asbestiform version, in your view you can determine
 14:09:18 **12** that a single fiber is asbestiform; is that your
 14:09:21 **13** view?
 14:09:21 **14 A. A single fiber in one sample, yes, if it's**
 14:09:25 **15 fibrous, it's asbestiform. It does not meet the**
 14:09:27 **16 definition of a geological grab sample because the**
 14:09:31 **17 definition of asbestiform provides you all these**
 14:09:34 **18 different tests that are not applicable and don't**
 14:09:37 **19 give you any measurement ability to measure high**
 14:09:42 **20 tensile strength. Don't even tell you what high**
 14:09:45 **21 tensile strength is. You can't measure flexibility.**
 14:09:47 **22 You can't measure -- you can't measure any of that**
 14:09:53 **23 with any of the protocols. And these are fibers that**
 14:09:59 **24 are regulated asbestos fibers. My definition is**
 14:10:03 **25 they're all asbestiform. Others can argue with that.**
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14:10:12 **1** MR. EWALD: Promise to not spend too much
 14:10:14 **2** time on 22262-1 today, but I have just a couple
 14:10:18 **3** of questions. So as Exhibit 9, we're going to
 14:10:35 **4** mark ISO 22262-1.
 14:10:43 **5** (Defendant's Exhibit 9 was marked for
 14:10:55 **6** identification.)
 14:10:55 **7 Q.** (By Mr. Ewald) Doctor, this ISO 22262-1
 14:10:59 **8** is one of the methods you used in connection with
 14:11:02 **9** your November 14th, 2018, report; is that correct?
 14:11:08 **10 A. That is correct.**
 14:11:21 **11 Q.** You've been asked for --
 14:11:25 **12 A. 2.8?**
 14:11:26 **13 Q.** -- about 2.8. I'm not going to go back
 14:11:29 **14** there, at least immediately.
 14:11:31 **15** Am I correct that you referred to 2.8 on
 14:11:36 **16** page 2, definition of asbestiform as, quote, a
 14:11:43 **17** general definition; is that right?
 14:11:44 **18 A. That's correct.**
 14:11:46 **19 Q.** Okay. Let's go to page 23, and that is
 14:11:53 **20** carryover of 7.2.3.7.1, morphology. Let me know when
 14:11:59 **21** you get to 23.
 14:12:03 **22 A. I have it.**
 14:12:04 **23 Q.** Okay. Do you agree with me that ISO
 14:12:18 **24** 22262-1 distinguishes in this section between
 14:12:26 **25** asbestiform amphibole fibers and nonasbestiform
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14:12:30 **1** amphibole fibers?
 14:12:43 **2** MR. HORN: Can I have that question read
 14:12:46 **3** back? I was actually reading the protocol.
 14:13:09 **4** (Record was read by the court reporter.)
 14:13:09 **5** MR. HORN: Thank you.
 14:13:09 **6** THE WITNESS: Yes. This is intended as
 14:13:11 **7** guidance for analysts to discriminate -- from
 14:13:15 **8** the note on page 23 -- between nonasbestiform
 14:13:23 **9** and asbestiform amphibole populations. It is
 14:13:27 **10** not intended to override the definition of
 14:13:29 **11** asbestos as presented in 2.9 nor override any
 14:13:33 **12** national regulation.
 14:13:35 **13** So according to that, it is guidance to
 14:13:38 **14** the analyst.
 14:13:41 **15 Q.** (By Mr. Ewald) Okay. You can keep that
 14:14:02 **16** close by. We'll come back to it maybe.
 14:14:05 **17** I want to talk a little bit about
 14:14:17 **18** Mr. Poye.
 14:14:17 **19 A. Sure.**
 14:14:25 **20** MR. EWALD: Let's mark this as Exhibit 10.
 14:14:25 **21** (Defendant's Exhibit 10 was marked for
 14:14:43 **22** identification.)
 14:14:43 **23 Q.** (By Mr. Ewald) Doctor, as Exhibit 10,
 14:14:52 **24** handing you what is -- it's got a couple pages at the
 14:14:58 **25** beginning. Can you describe what those first couple
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14:15:01 **1** pages are?

14:15:01 **2 A. The chain of custodies for the samples we**

14:15:07 **3 sent him. I think there's 50 -- 79. These are the**

14:15:22 **4 chain of custodies for the 79 samples we sent him.**

14:15:25 **5 Also in the chain of custody is the key because we**

14:15:28 **6 sent them as blind, meaning we generated a number**

14:15:32 **7 68708, 1 through 79, and then the corresponding MDL**

14:15:40 **8 samples that we have. And then on the back -- on**

14:15:50 **9 another page has his numbers, then the actual MDL**

14:15:55 **10 samples that were sent.**

14:16:07 **11 Q. Am I correct after that you have**

14:16:12 **12 Mr. Poye's July 18th, 2018, report to you on the MAS**

14:16:17 **13 split of historic talc samples?**

14:16:20 **14 A. Correct.**

14:16:21 **15 Q. And have you seen this before?**

14:16:23 **16 A. Of course.**

14:16:26 **17 Q. Just making sure.**

14:16:29 **18 Now, when was the last time you spoke to**

14:16:39 **19 Mr. Poye about the testing reflected in his**

14:16:46 **20 July 18th, 2018, report?**

14:16:48 **21 A. Some months. I don't remember the last**

14:16:50 **22 time I talked to him about it.**

14:16:52 **23 Q. Do you recall whether or not it was before**

14:16:57 **24 or after he sent you this work?**

14:17:03 **25 A. No, I don't recall.**

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14:17:05 **1 Q. What do you recall about that**

14:17:08 **2 communication?**

14:17:09 **3 A. I don't recall any of it.**

14:17:11 **4 Q. Was it by phone? In person? Email?**

14:17:14 **5 A. I'm sure it would have been by phone. I**

14:17:19 **6 know I talked to him about sending the samples. Have**

14:17:22 **7 I talked to him about the results or not, I don't**

14:17:24 **8 have any recollection of that.**

14:17:26 **9 Q. Have you communicated with anyone other**

14:17:32 **10 than Mr. Poye at his lab, J3, about the testing that**

14:17:39 **11 is reflected in his July 18th report?**

14:17:42 **12 A. Not that I recall.**

14:17:45 **13 Q. There was some confusion, I believe,**

14:17:54 **14 during --**

14:17:55 **15 A. Chain of custody of the spike samples?**

14:18:01 **16 Q. Actually, that wasn't where I was going**

14:18:06 **17 but --**

14:18:06 **18 A. Everybody seems to have confusion on that,**

14:18:10 **19 and I still have to get the chain of custody to**

14:18:12 **20 explain how the numbers work on the spike samples.**

14:18:15 **21 But I shouldn't cut you off and generate more**

14:18:18 **22 questions.**

14:18:18 **23 Q. Exactly. I was going there next.**

14:18:23 **24 Actually, it seemed to be unclear during**

14:18:27 **25 the Leavitt depositions about what method Lee Poye**

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14:18:36 **1** used with respect to PLM, and I would like you to

14:18:41 **2** turn to --

14:18:42 **3 A. Used the ISO method.**

14:18:44 **4 Q. Okay. It's your understanding that he**

14:18:46 **5** used the ISO 22262-1 method?

14:18:54 **6 A. Correct. And the confusion came in where**

14:18:59 **7 the original XRD samples I sent him for those 30**

14:19:01 **8 samples, I believe that was the R-93 method. I just**

14:19:03 **9 got them mixed up in my head. I could have just**

14:19:03 **10 looked at the document and stated it correctly.**

14:19:15 **11 Q. Do you have an understanding as to why J3**

14:19:25 **12** used ISO 22262-1 as opposed to R-93 for the PLM work?

14:19:31 **13 A. I'll have to let Lee Poye speak to that.**

14:19:36 **14 It looks like to me just standardizing the protocol**

14:19:39 **15 since there is both an ISO PLM for talc as well as an**

14:19:43 **16 ISO TEM for talc.**

14:19:48 **17 Q. So I asked Lee Poye that.**

14:19:50 **18 A. What did he say?**

14:19:52 **19 Q. Mr. Poye said that his recollection is**

14:19:54 **20** that you told him to use ISO 22262-1. Do you have

14:19:59 **21** any recollection of that?

14:20:00 **22 A. It's possible. I just don't recall that.**

14:20:03 **23 Q. Do you see on -- it's the cover letter to**

14:20:08 **24** the report on top of page 2, at the top, first full

14:20:18 **25** sentence, J3 was directed to analyze the talcum

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14:20:21 **1** powder samples for the presence and percentage of

14:20:24 **2** regulated asbestos utilizing the following

14:20:27 **3** appropriate methods, and one of the methods there

14:20:30 **4** that's listed is ISO 22262-1; do you see that?

14:20:38 **5 A. I do see that. Now, if that's Mr. Poye's**

14:20:43 **6 recollection, I have no reason to dispute that.**

14:20:45 **7 Q. I take it, then, you have not reviewed**

14:20:48 **8** Mr. Poye's deposition in the Lovett case relating to

14:20:52 **9** testing of the samples reflected in the July 18th

14:20:56 **10** report?

14:20:56 **11 A. I have not.**

14:21:13 **12 Q. Do you have any knowledge about J3's**

14:21:20 **13** testing of the MDL samples beyond what is reflected

14:21:28 **14** in this July 18, 2018, report?

14:21:33 **15 A. I don't think so.**

14:21:41 **16 Q. Let me toggle between a couple of reports**

14:21:46 **17** now. I'll keep the Lee Poye one close by, but if

14:21:49 **18** you'll look at your November report, on page 16

14:22:03 **19** there's a reference to the comparing of -- well, let

14:22:11 **20** me take a step back.

14:22:12 **21** Can you please explain what you asked Lee

14:22:28 **22** Poye's lab, J3, to do with respect to the 79 samples

14:22:37 **23** that you sent them.

14:22:38 **24 A. Well, it looks like I asked them to**

14:22:41 **25 analyze it by the ISO 22262-1 protocol for PLM and**

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14:22:48 **1 XRD.**

14:22:52 **2 Q.** And then, at least as of the time of this

14:22:55 **3** report, this report being the November 14th report,

14:23:04 **4** J3 and MAS had analyzed 22 of same samples by ISO PLM

14:23:27 **5** that MAS had; is that correct?

14:23:28 **6 A. That's correct.**

14:23:29 **7 Q.** And the statement here at the bottom that

14:23:35 **8** J3 did not find any asbestos detected in those 22

14:23:42 **9** samples, while MAS found asbestos in eight of the 22

14:23:47 **10** samples using the ISO PLM method, is that the correct

14:23:51 **11** statement?

14:23:52 **12 A. That is a correct statement.**

14:23:53 **13 Q.** And then it goes on to state, These

14:23:56 **14** different results between the two labs will require

14:23:59 **15** further investigation to understand the reason for

14:24:01 **16** these differences.

14:24:02 **17** Did I read that correctly?

14:24:04 **18 A. You did.**

14:24:04 **19 Q.** As of day 3 of Leavitt, I think this was

14:24:09 **20** asked -- maybe it was day two --

14:24:12 **21 A. Or day one --**

14:24:14 **22 Q.** There had not been any steps taken at that

14:24:17 **23** point in time on this question of investigating the

14:24:19 **24** different results. Have there since been any efforts

14:24:24 **25** by MAS or efforts you're aware of from J3 to

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14:24:28 **1** investigate the different results reached on those 22

14:24:32 **2** PLM samples?

14:24:32 **3 A. We haven't done anything further. May**

14:24:36 **4 develop some standards at the lower detection limit**

14:24:39 **5 and have a blind round-robin between the two analysts**

14:24:43 **6 to see how consistent they are.**

14:24:45 **7 But other than that, I don't have any**

14:24:48 **8 other information that I can share for the reason for**

14:24:52 **9 the eight positives versus the eight negatives by J3.**

14:24:57 **10 Q.** So as you sit here today, do you have any

14:25:01 **11** basis to understand why two labs using the same

14:25:09 **12** method and the same samples reached different results

14:25:12 **13** in eight cases?

14:25:14 **14 A. For eight samples? No.**

14:25:21 **15 Q.** Do you know who did the PLM work at J3?

14:25:27 **16 A. Other than it's their PLM analyst, I can't**

14:25:38 **17 remember that individual's name right at the moment.**

14:25:41 **18 Q.** Without remembering his or her name, do

14:25:48 **19** you know if you met that person before?

14:25:49 **20 A. I've been to their lab. I don't know if I**

14:25:56 **21 met him or not.**

14:25:57 **22 Q.** It's your recollection it's a he, not a

14:26:00 **23** she?

14:26:01 **24 A. To my recollection it's a he.**

14:26:05 **25 Q.** I think we're done with Poye, at least for

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14:26:27 **1** now, and can set that aside.

14:26:34 **2** Going back to your November report, what

14:26:40 **3** role did Dr. Rigler play in this report?

14:26:44 **4 A. He helped edit and review the data so that**

14:26:48 **5 he was one of the QC people.**

14:27:18 **6 Q.** Why did you ask J3 to perform the ISO PLM

14:27:25 **7** on the same samples that MAS was testing with ISO

14:27:30 **8** PLM?

14:27:30 **9** MR. HORN: Object to form.

14:27:33 **10** THE WITNESS: They had initially done a

14:27:37 **11** number of the ISO PLM. We had sent off the

14:27:41 **12** samples for the original 30 and, just to be

14:27:43 **13 consistent, wanted to see the difference between**

14:27:47 **14 the two labs that used that method.** And of

14:27:54 **15** course the XRD, we don't have an XRD. So it

14:28:00 **16** just was interesting to me.

14:28:01 **17 Q.** (By Mr. Ewald) Trying to get a sense of,

14:28:36 **18** essentially, the workflow for the November 14th

14:28:39 **19** testing -- with respect to the testing reflected in

14:28:45 **20** the November 14th report at MAS, and there are a

14:28:50 **21** number of different test methods that are described

14:28:52 **22** there that MAS conducted. Do you have a sense of how

14:28:56 **23** that played out, whether it was all done in the same

14:28:59 **24** order with respect to a particular sample? Can you

14:29:03 **25** give some explanation of that?

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14:29:05 **1 A. When you say a particular order --**

14:29:07 **2 Q.** So, for example, you have one of the 56

14:29:15 **3** samples there that MAS tested by ISO PLM, Blount PLM,

14:29:23 **4** and TEM; right?

14:29:26 **5 A. Correct.**

14:29:27 **6 Q.** For each sample was the analysis done in a

14:29:32 **7** particular order with respect to the type of testing,

14:29:38 **8** or was it just done in whatever order it happened?

14:29:43 **9 A. Well, there's two different groups that**

14:29:45 **10 are doing the testing, so you have the PLM group and**

14:29:50 **11 then the TEM group. Typically we don't -- we're not**

14:29:57 **12 trying to do one before the other because they're two**

14:30:00 **13 separate groups. And irrespective of the ISO**

14:30:05 **14 22262-1, which says you should do the PLM first,**

14:30:09 **15 somebody's already asked that because some of the**

14:30:11 **16 PLMs -- was it you?**

14:30:13 **17 Q.** Yeah, that would be me.

14:30:15 **18 A. It was a good question. A good question**

14:30:17 **19 in that I believe the reason for that is instead of**

14:30:20 **20 fully characterizing these cosmetic talcs like we're**

14:30:25 **21 doing, if you find it by PLM and it meets the**

14:30:29 **22 definition, probably what I believe -- I should email**

14:30:35 **23 Dr. Chatfield -- Eric -- one of these days -- is that**

14:30:40 **24 there's really no need to go any further if you're**

14:30:45 **25 just trying to determine if there's asbestos or not**

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14:30:47 **1 present.**

14:30:47 **2 There's four TEM analysts. Whatever they**

14:30:50 **3 seem to be working on, some will take samples -- say**

14:30:54 **4 we have 22 samples. Some may be working from 22 on**

14:30:59 **5 down, some may be grabbing in the middle. It's just**

14:31:02 **6 whoever's free and what's available, what's been**

14:31:05 **7 prepared. At the end of the day then we're just**

14:31:10 **8 trying to put them all together. However, I'll give**

14:31:15 **9 you credit now trying to get the PLM analysts to get**

14:31:19 **10 ahead of the TEM guys.**

14:31:22 **11 Q.** Have you communicated with Dr. Chatfield

14:31:25 **12** in any way about his -- not his -- about ISO 22262-1?

14:31:33 **13 A.** Well, to give Dr. -- Eric credit, it's

14:31:37 **14 basically his.**

14:31:38 **15 Q.** The answer to my question is yes or no?

14:31:40 **16 A.** No, I have not talked to Eric. I haven't

14:31:43 **17 seen him in a while. He is a world of knowledge.**

14:31:51 **18 Q.** Do you agree that it's common for

14:31:57 **19** independent labs to design standard operating

14:32:00 **20** procedures for companies analyzing asbestos in their

14:32:03 **21** products?

14:32:05 **22 A.** Is it standard operating procedure --

14:32:10 **23** MR. HORN: Let me just object to form.

14:32:12 **24** It's vague and ambiguous.

14:32:13 **25** THE WITNESS: And I'm just trying to

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14:32:15 **1** clarify, so that if a company comes and says

14:32:17 **2** develop a protocol for me, I mean, we've done

14:32:20 **3** that once. We did it for Federal-Mogul for when

14:32:24 **4** they came into an accessory mineral tremolite

14:32:27 **5** issue with their brake shoes.

14:32:35 **6** So I can't speak for other labs. What I

14:32:38 **7** know is that we had developed a protocol for

14:32:41 **8** them, and I guess they -- if you go on Google,

14:32:46 **9** you can get it.

14:32:46 **10 Q.** (By Mr. Ewald) And did that relate to

14:32:51 **11** analyzing raw wollastonite?

14:32:56 **12 A.** It did.

14:32:57 **13 Q.** And that was with PLM?

14:32:59 **14 A.** It was a combination, but primarily PLM

14:33:04 **15** after we went through a concentration method.

14:33:07 **16** Because of the concentration that was being found in

14:33:09 **17** there, you only needed PLM.

14:33:14 **18 Q.** And that one, the SOP used R-93?

14:33:20 **19 A.** Yes, I believe so. I haven't seen that

14:33:24 **20** SOP in a while, and you're probably looking at it.

14:33:33 **21 Q.** So you don't have a sense one way or the

14:33:53 **22** other, apart from your own lab, about how often that

14:33:56 **23** happens in the industry with respect to other labs?

14:33:59 **24 A.** No. Obviously, I'm aware that the McCrone

14:34:04 **25** TEM method is identical to Johnson & Johnson's 7024.

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14:34:12 **1 I'm assuming McCrone is the ones that developed a**

14:34:15 **2 method that was adopted by Johnson & Johnson.**

14:34:23 **3 Outside of that, I really don't have a**

14:34:25 **4 sense on what other commercial laboratories have done**

14:34:27 **5 for clients.**

14:34:28 **6 Q.** And you don't hold the opinion that the

14:34:31 **7** McCrone and Millette method was designed as a rigged

14:34:37 **8** test, do you?

14:34:37 **9 A.** As a what?

14:34:38 **10 Q.** As a rigged test.

14:34:39 **11 A.** I don't know what rigged means. I don't

14:35:12 **12** know what McCrone was thinking.

14:35:17 **13 Only issue I have with all those methods**

14:35:19 **14** is that there seems to be a lack of understanding on

14:35:22 **15** just how much asbestos -- regulated asbestos meeting

14:35:29 **16** definitions has to be in the sample before you call

14:35:32 **17** it a positive sample.

14:35:35 **18 And as I've progressed through this work,**

14:35:37 **19** the weight percents -- and that's the other problem I

14:35:41 **20** have, is the theoretical weight percent detection

14:35:47 **21** limits.

14:35:48 **22 Where the disconnect on that is our**

14:35:52 **23** detection limit is based on one small fiber, and it's

14:35:58 **24** like looking at the FDA 2010 data from AMA, and I

14:36:04 **25** think they gave a detection limit of 2.0 times 10 to

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14:36:09 **1 the minus 6 weight percent because they're detecting**

14:36:13 **2 one fiber. Well, you have to understand in order to**

14:36:16 **3** detect that one fiber for the TEM method that AMA was

14:36:20 **4** looking at, you have to have something like 12

14:36:23 **5** million fibers before, statistically, you may run

14:36:26 **6** across one fiber.

14:36:28 **7 So the method is the method. I'll let**

14:36:31 **8** others decide if it's, quote, rigged or not. But you

14:36:36 **9** can't use that method and say there's nothing in the

14:36:45 **10** talc. That's my only issue with it.

14:36:47 **11 Q.** Have you reviewed any reports from

14:36:49 **12** Dr. Sanchez that he recently released relating to the

14:36:53 **13** MDL samples?

14:36:54 **14 A.** Yes, I have.

14:36:55 **15 Q.** Do you recall which reports you reviewed?

14:36:59 **16 A.** I understand he has them all, but I think

14:37:02 **17** the first ten or so, which were the ten samples from

14:37:06 **18** Vermont where he says all the anthophyllite that --

14:37:15 **19** well, the anthophyllite series that we identified is

14:37:19 **20** all nonasbestos -- is all cleavage fragment

14:37:26 **21** cummingtonite.

14:37:27 **22 Q.** What's your response to that?

14:37:29 **23 A.** Well, my comments are, one, it's unclear

14:38:08 **24** to me why he used electron backscatter detection for

14:38:20 **25** determining the crystalline structure. What he did

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15:57:24 **1 asbestos, and that would be more accurate for the**
 15:57:26 **2 detection limit for weight percent.**
 15:57:28 **3 I don't mean to be thinking out loud, but**
 15:57:30 **4 I think in the supplement report that's something we**
 15:57:33 **5 can do. That way you're not picking something. You**
 15:57:39 **6 say, okay, for this I measured 60 or 70 asbestos**
 15:57:43 **7 structures and it came from the Vermont mine and**
 15:57:46 **8 here's the average length and width, so I'm going to**
 15:57:49 **9 use that for the analytical sensitivity.**
 15:57:52 **10 I don't mean to come up with an idea**
 15:57:54 **11 sitting here.**
 15:57:56 **12 MR. HORN: Just created more work.**
 15:57:58 **13 THE WITNESS: It's the nerd in me, I**
 15:58:01 **14 guess.**
 15:58:02 **15 Q. (By Mr. Horn) And just so it's clear for**
 15:58:04 **16 the record, the analytical sensitivity, when you use**
 15:58:09 **17 it with a weight percentage, is that just dictated by**
 15:58:12 **18 the protocol that you're using? Or when would you**
 15:58:15 **19 use that versus something else?**
 15:58:16 **20 A. Well, most of these protocols call for a**
 15:58:18 **21 weight percent. But in order to have a weight**
 15:58:20 **22 percent, you have to count all the number of fibers**
 15:58:22 **23 and structures that are in there. So it's literally**
 15:58:26 **24 how you report it.**
 15:58:27 **25 Nothing changes how you collect the data.**

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15:58:30 **1 So any of these TEM count sheets, you would take the**
 15:58:34 **2 average length and width of everything in there for**
 15:58:38 **3 these weight percents that we have reported.**
 15:58:41 **4 For example, Table 4, we found a**
 15:58:45 **5 concentration of 53,000 asbestos structures per gram.**
 15:58:51 **6 That number of fibers that made that 53,000 is the**
 15:58:55 **7 same number of fibers that you would do the exact**
 15:58:58 **8 same calculations. Instead of fibers per gram, it's**
 15:59:03 **9 going through and measuring every length, every**
 15:59:08 **10 width, and typically they end up in picograms of some**
 15:59:13 **11 amount, and that's added all up, and then you go**
 15:59:16 **12 through the exact same calculations that you go**
 15:59:18 **13 through to determine how many fibers or bundles per**
 15:59:21 **14 gram. It's just the way they express the data.**
 15:59:24 **15 In my opinion, it is better to have fibers**
 15:59:27 **16 and bundles per gram, because if you measure an**
 15:59:31 **17 exposure, it's always fibers per cubic centimeter of**
 15:59:37 **18 air. It's not weight percent of a cubic centimeter**
 15:59:40 **19 of air. That's been shown to be completely**
 15:59:43 **20 inaccurate, and it doesn't tell you anything about**
 15:59:45 **21 your potential exposure.**
 15:59:46 **22 That's why I believe the fibers per gram**
 15:59:50 **23 or fiber bundles per gram is important, because it**
 15:59:53 **24 gives you some idea of the potential for exposure**
 15:59:56 **25 when measuring fibers per cc.**

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15:59:57 **1 MR. HORN: That's all I have.**
 16:00:02 **2 MR. EWALD: I'm just going to put on the**
 16:00:03 **3 record, Ethan, I just haven't seen the letter or**
 16:00:07 **4 the back and forth you referenced about Leavitt.**
 16:00:09 **5 I don't need to reserve any rights, but I'm not**
 16:00:12 **6 familiar with it so I just wanted to make note**
 16:00:15 **7 of it in case people take issue with it.**
 16:00:20 **8 MR. HORN: We sent a link out that**
 16:00:23 **9 contained a lot of materials, and it's tucked in**
 16:00:26 **10 there.**
 16:00:29 **11 MR. EWALD: All right.**
 16:00:30 **12 MR. HORN: Before we go, let me just email**
 16:00:33 **13 it to both of you.**
 16:07:06 **14 (Deposition concluded at 4:00 p.m.)**
15 (Pursuant to Rule 30(e) of the Federal
16 Rules of Civil Procedure and/or OCGA 9-11-30(e),
17 signature of the witness has been reserved.)
18 (Original transcript sent to McCarter &
19 English, LLP.)
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1 C E R T I F I C A T E
2
3 STATE OF GEORGIA:
4 COUNTY OF GWINNETT:
5
6 I hereby certify that the foregoing
7 transcript was taken down, as stated in the
8 caption, and the questions and answers thereto
9 were reduced to typewriting under my direction;
10 that the foregoing pages 1 through 131 represent
11 a true, complete, and correct transcript of the
12 evidence given upon said hearing, and I further
13 certify that I am not of kin or counsel to the
14 parties in the case; am not in the regular
15 employ of counsel for any of said parties; nor
16 am I in anywise interested in the result of said
17 case.
18 This, the 10th day of January 2019.
19
20
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25

DEBRA R. LUTHER, B-881
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DEPOSITION OF WILLIAM E. LONGO, PhD /DRL

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If supplemental or additional pages are necessary, please furnish same in typewriting annexed to this deposition.

WILLIAM E. LONGO, PhD

Sworn to and subscribed before me,

This, the ____ day of _____, 20____.

Notary Public
My commission expires: _____

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DEPOSITION OF WILLIAM E. LONGO, PhD /DRL

I do hereby certify that I have read all questions propounded to me and all answers given by me on the 7th day of January 2019, taken before Debra R. Luther, and that:

- ____ 1) There are no changes noted.
____ 2) The following changes are noted:

Pursuant to Rule 30(e) of the Federal Rules of Civil Procedure and/or the Official Code of Georgia Annotated 9-11-30(e), both of which read in part: Any changes in form or substance which you desire to make shall be entered upon the deposition...with a statement of the reasons given...for making them. Accordingly, to assist you in effecting corrections, please use the form below:

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